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**Efficacy and Safety of Asfotase Alfa in Infants and Young Children With
Hypophosphatasia: a Phase 2 Open-Label Study**

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Short title: Asfotase alfa in infants/young children with HPP

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Conflict of interest disclosures

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70 M has been an investigator and has received grant funding and/or research support
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73

74 [word count limit: 250 words (structured); current count: 250]

75 **ABSTRACT**

76 **Context:** Long-term data on enzyme replacement treatment of hypophosphatasia
77 (HPP) are limited.

78 **Objective:** To evaluate efficacy and safety of asfotase alfa in patients aged ≤ 5 years
79 with HPP followed for up to 6 years.

80 **Design:** Phase 2 open-label study (July 2010–September 2016).

81 **Setting:** 22 sites, 12 countries.

82 **Participants:** Sixty-nine patients (median [range] age : 16.0 [0.02–72] mo) with severe
83 HPP and sign/symptom-onset before age 6 months.

84 **Intervention:** Asfotase alfa 2 mg/kg 3 times/wk or 1 mg/kg 6 times/wk subcutaneously.

85 **Main Outcome Measures:** Primary efficacy measure: Radiographic Global Impression
86 of Change (RGI-C) score (-3 [severe worsening] to $+3$ [complete/near-complete
87 healing]). Additional outcome measures: respiratory status, growth, safety. Post hoc
88 analysis: characteristics of radiographic “responders” vs. “nonresponders” at Year 1
89 (RGI-C: $\geq +2$ vs. $< +2$).

90 **Results:** During median (min, max) 2.3 (0.02, 5.8) years of treatment, RGI-C scores
91 improved significantly at Month 6 ($+2.0$ [-1.7 , $+3.0$]), Year 1 ($+2.0$ [-2.3 , $+3.0$]), and Last
92 Assessment ($+2.3$ [-2.7 , $+3.0$]; $P < .0001$ all). Of 24 patients requiring respiratory support
93 at Baseline, 11 (46%) no longer needed support. Height/weight Z-scores generally
94 increased. Nine patients died (13%). All patients experienced ≥ 1 adverse event; pyrexia
95 was most common. Compared with “responders” ($n=50$ [72%]), “nonresponders” ($n=19$

[28%]) had more severe disease at Baseline and a higher rate of neutralizing antibodies at Last Assessment.

Conclusions: Most infants/young children treated with asfotase alfa showed early radiographic and clinical improvement that was sustained up to 6 years; radiographic “nonresponders” had more severe underlying disease and more frequent neutralizing antibodies at Last Assessment.

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Précis

Most infants and young children with hypophosphatasia treated with asfotase alfa showed improved skeletal manifestations, respiratory function, and growth within 1 year, maintained up to 6 years.

109 [word count limit: none; current count: 4793]

110 **INTRODUCTION**

111 Hypophosphatasia (HPP) is the rare, inherited, systemic, metabolic disease
112 characterized by low activity of the tissue-nonspecific isoenzyme of alkaline
113 phosphatase (TNSALP), which leads to extracellular accumulation of its substrates,
114 mainly inorganic pyrophosphate (PPi) and pyridoxal 5'-phosphate (PLP) (1-4).
115 Increased extracellular levels of PPi inhibit bone mineralization and lead to impaired
116 skeletal mineralization in affected patients and additional rickets-like deformities in
117 infants and children (2,4). Reduced dephosphorylation of PLP, the circulating form of
118 vitamin B6, by TNSALP has been associated with vitamin B6-responsive seizures in
119 infants with HPP (3,5).

120

121 Clinical presentation of HPP varies with age at onset, from in utero to adulthood (2,6).
122 Characteristic signs, symptoms, and complications of perinatal and infantile HPP that
123 are potentially life-threatening include respiratory failure, vitamin B6-responsive
124 seizures, chest deformity, and craniosynostosis; other manifestations include severe
125 hypercalcemia, nephrocalcinosis, poor growth, osteomalacia, and bowing of the long
126 bones (2,5,7-11). Historically, patients with perinatal and infantile HPP have 58% to
127 100% mortality during the first year of life (12-14). The most common cause of death
128 among infants with HPP is respiratory failure secondary chest deformity and pulmonary
129 hypoplasia (12,15).

130

Asfotase alfa (Strensiq®; Alexion Pharmaceuticals, Inc., Boston, MA, USA) is a human recombinant TNSALP enzyme replacement therapy approved for patients with pediatric-onset HPP (16). In an open-label study of 11 infants and young children (aged ≤ 3 y) with life-threatening HPP, treatment with asfotase alfa for up to 7 years improved HPP-related skeletal abnormalities seen on radiograph, respiratory function, growth, and cognitive and motor function (17,18). Here we report the long-term safety and efficacy of asfotase alfa in the largest study to date of infants and children aged ≤ 5 years with manifestations of HPP before age 6 months.

MATERIALS AND METHODS

Patients

Children aged ≤ 5 years with signs or symptoms of HPP before age 6 months were eligible for enrollment if they had a documented diagnosis of HPP. Diagnosis of HPP required the following: total serum alkaline phosphatase (ALP) activity below the lower limit of normal for age, plasma PLP above the upper limit of normal (unless the patient was receiving pyridoxine for seizures), radiographic evidence of HPP (flared and frayed metaphyses, widened growth plates, areas of radiolucency or sclerosis, or severe, generalized osteopenia), and ≥ 2 HPP-related findings (history or presence of nontraumatic postnatal fracture and/or delayed fracture healing, nephrocalcinosis or history of elevated serum calcium, functional craniosynostosis, respiratory compromise or rachitic chest deformity, vitamin B6-responsive seizures, or failure to thrive). Exclusion criteria were serum calcium or phosphate levels below the normal range, serum 25(OH) vitamin D levels < 20 ng/mL, current evidence of a treatable form of

rickets, prior treatment with bisphosphonates, investigational drug treatment within 1 month, or current enrollment in any other study involving a new drug, device, or treatment for HPP.

The study complied with the Declaration of Helsinki and International Conference on Harmonisation Guideline for Good Clinical Practice and with national, state, and local laws of pertinent regulatory authorities. The protocol was approved by each site's institutional review board/independent ethics committee, and written informed consent was obtained for all patients from a parent(s) or guardian(s).

Study Design

In this open-label, multicenter, single-arm, multinational study (ClinicalTrials.gov NCT01176266; EUDRACT 2010-019850-42), eligible patients received a total subcutaneous dose of 6 mg/kg/wk of asfotase alfa administered as 1 mg/kg 6 times per week or 2 mg/kg 3 times per week (maximum volume: 1 mL asfotase alfa per injection). Dose adjustments were allowed at the investigator's discretion to account for changes in body weight or in consultation with the medical monitor for safety concerns or lack of efficacy. The maximum dose permitted was 40 mg per injection or 9 mg/kg/wk in Australia, France, Germany, Italy, Saudi Arabia, Spain, and the United Kingdom per protocol amendment; no dose restrictions were applied in Canada, Japan, Russia, Turkey, or the United States. The initial dose of asfotase alfa was administered at the study site during the Baseline visit; post-Baseline injections could be administered at home by a parent, legal guardian, or designee after adequate training. With each

injection, the designated individual was required to complete a worksheet regarding the patient's health condition, any new medications, and details of the injection. Study visits were scheduled at Weeks 3 and 6, Months 3, 6, 9, 12, 15, 18, and 24, and every 6 months thereafter until the end-of-study assessment (Month 48 in patients enrolled in the United Kingdom and final every-6-month assessment in other countries). Patients were enrolled starting July 22, 2010, and the last patient completed the study September 26, 2016.

Outcomes Measures

Primary efficacy measure

The primary efficacy measure was improvement of HPP-related skeletal manifestations at Week 24 (Month 6) and Week 48 (Year 1) of treatment as measured on the Radiographic Global Impression of Change (RGI-C) scale (19). The RGI-C (19) is a validated 7-point scale that assesses changes from Baseline in HPP-related skeletal abnormalities: -3=severe worsening, -2=moderate worsening, -1=minimal worsening, 0=no change, +1=minimal healing, +2=substantial healing, and +3=complete or near complete healing. Radiographs of the chest, bilateral wrists, and bilateral knees were reviewed by 3 independent pediatric radiologists, and comparisons with Baseline were scored. The mean RGI-C score for each patient at each time point was calculated from available scores. The radiologists were blinded to post-Baseline timepoints and all other patient information.

Secondary efficacy measures

Skeletal manifestations of HPP over time: RGI-C scores and change from Baseline in Rickets Severity Scale (RSS) (20) scores were assessed at all study visits starting at Month 3. The RSS (20) is a 10-point scale (0=absence of metaphyseal cupping and fraying [both characteristic of rickets] to 10=severe rickets; maximum of 4 points for the wrists and 6 points for the knees) originally developed to assess the severity of nutritional rickets in the wrists and knees. Radiographs for determination of RSS score were read by a single independent rater who developed the RSS (Tom D. Thacher, MD). The percentage of “responders” (individual mean RGI-C score: $\geq +2$) at each study visit was also determined.

Respiratory status: Respiratory status (including use and type of support) was assessed at Screening, Baseline, and all subsequent study visits.

Growth: Length/height, weight, and head circumference were recorded during physical examinations at required study visits to assess changes in growth. Length/height and weight Z-scores were assigned based on the Centers for Disease Control and Prevention growth charts for age- and sex-matched healthy infants and children (21). Head circumference Z-scores were calculated using World Health Organization formulae (22).

Ventilator-free and overall survival: Ventilator-free survival was assessed with the occurrence of death and ventilatory support (continuous positive airway pressure

[CPAP], bilevel or biphasic positive airway pressure, or mechanical ventilation [invasive ventilation via endotracheal intubation or tracheostomy]). Supplemental oxygen was considered respiratory but not ventilatory support. Survival was monitored throughout the study.

Other measures

Blood samples were collected to assess serum ALP activity, plasma PPi and PLP, and serum parathyroid hormone (PTH) concentrations at required study visits after an overnight fast and before study drug administration.

ALPL gene mutation analysis for patients not previously tested was performed by Connective Tissue Gene Tests (Allentown, PA, USA).

Safety and tolerability

Safety was assessed by routine reporting of adverse events (AEs), which included serious AEs, injection site reactions (ISRs), and injection-associated reactions (IARs). ISRs were defined as treatment-emergent AEs (TEAEs) that were localized to the site of study drug administration, occurred at any time point after study drug initiation, and were assessed by the investigator as possibly, probably, or definitely related to study drug. IARs were defined as systemic signs, symptoms, or findings that occurred within 3 hours after study drug administration and were assessed by the investigator as possibly, probably, or definitely related to study drug. AEs of special interest included ectopic calcifications, lipodystrophy, craniosynostosis, and chronic hepatitis and were

based on clinical review of observed AEs. Additional safety assessments included physical examinations, clinical laboratory tests (including calcium and magnesium), anti-asfotase alfa antibody levels (PPD Laboratories, LLC, Richmond, VA, USA), fundoscopic eye examinations, and renal ultrasounds. The clinical significance of abnormal laboratory findings was judged by the investigator. Safety events reported after the study ended were not included.

“Responder” analysis

A post hoc analysis compared Baseline characteristics of “responders” by radiography (RGI-C score: $\geq +2$) with those of “nonresponders” (score: $< +2$) at Year 1 of treatment.

Statistical analysis

All efficacy and safety analyses were performed on the full analysis population (patients who received ≥ 1 dose of asfotase alfa). Some analyses were repeated on the per protocol population (patients who received any asfotase alfa and had no major protocol deviations that could influence treatment effect). In general, continuous variables were summarized descriptively (data reported herein are median [minimum, maximum] unless otherwise specified), and categorical variables were summarized by counts and percentages of patients.

For the primary efficacy analysis (RGI-C scores at Month 6 and Year 1), a nonparametric Wilcoxon signed-rank test was used to determine whether the median RGI-C scores at Month 6 and Year 1 differed from 0. Missing values were imputed

using last observation carried forward. Patients with no recorded post-Baseline values were assigned as having no change (score: 0).

Secondary efficacy analyses of RGI-C scores, percent of “responders,” and change from Baseline in RSS scores at each study visit were conducted in a manner similar to that used for the primary analysis; however, only observed data were used (no imputation). *P* values for length/height and weight Z-scores were calculated post hoc using the nonparametric Wilcoxon signed-rank test comparing median change to 0. Pharmacodynamic and safety assessments are summarized descriptively. Ventilator-free survival and overall survival time were assessed using Kaplan-Meier methodology.

For the post hoc “responder” analysis, *P* values were calculated using the exact Wilcoxon rank-sum test for continuous variables and the Fisher’s exact test for categorical variables. Missing values at Year 1 were imputed with last observation carried forward. Patients with no post-Baseline data were assigned as having no change (score: 0) and considered “nonresponders.”

RESULTS

Patients

In total, 69 patients were enrolled from 22 sites in 12 countries and were included in the full analysis population, and 57 were included in the per protocol population (**Figure 1**). Patients were excluded from the per protocol population if they did not meet entry criteria or violated entry criteria (n=9), if they deviated from study protocol procedures

(n=2), if study drug was administered incorrectly (n=2) or not at all (n=1), or if an assessment/procedure was not done (n=1) (**Figure 1**). The number of patients enrolled in each country was as follows: Australia (n=1), Canada (n=11), France (n=5), Germany (n=13), Italy (n=2), Japan (n=5), Saudi Arabia (n=1), Russia (n=1), Spain (n=1), Turkey (n=4), United Kingdom (n=4), and United States (n=21). Baseline demographic and clinical characteristics are summarized in **Table 1**.

Dosing

Nearly all patients (67/69 [97%]) started asfotase alfa at 6 mg/kg/wk, with 64 (96%) receiving 2 mg/kg 3 times per week and 3 (4%) receiving 1 mg/kg 6 times per week. One patient started at 2 mg/kg 7 times per week and another started at 3 mg/kg 3 times per week. Doses were increased or decreased to 3–28 mg/kg/wk for 17/69 (25%) patients to account for changes in body weight, to enhance the likelihood of a clinical response, or because of AEs and administration issues (volume and number of injections).

Overall, median treatment duration was 2.3 (0.02, 5.8) years. Of the 69 patients in the full analysis population, 3 (4%) received treatment for <3 months and 14 (20%) for ≥36 months.

Primary efficacy measure

At Month 6 of treatment, the median (min, max) RGI-C score indicated significant improvement (+2.0 [−1.7, +3.0]; $P<.0001$; n=69); most patients (40/69 [58%]) were

considered “responders,” and 6 (9%) achieved a score of +3, indicating “complete or near complete healing” of HPP-related skeletal manifestations. Results observed at Month 6 were consistent with those at Year 1 (+2.0 [−2.3, +3.0]; $P<.0001$; $n=69$); 50/69 (72%) patients were considered “responders,” of which 4 (6%) achieved a score of +3. Results were similar in the per protocol population (data not shown).

Secondary efficacy measures

Skeletal manifestations of HPP over time: Preliminary patient-level radiographic outcomes have been published (14). Significant ($P<.05$) improvements in RGI-C score were observed at Months 3 and 6, Years 1, 2, 3, 4, and 5, and Last Assessment (**Figure 2**). The proportion of patients classified as “responders” (RGI-C score $\geq +2$) increased during the study, from 36% (24/66 patients) at Month 3 to 73% (49/67 patients) at Last Assessment. Consistent with RGI-C scores, RSS scores improved significantly ($P<.05$) from Baseline at Months 3 and 6, Years 1, 2, 3, 4, and 5, and Last Assessment (**Figure 3**). Results were similar in the per protocol population (data not shown).

Respiratory status: Of the 45/69 (65%) patients who did not require respiratory support at Baseline, 38 (84%) lived without support during the study and 43 (96%) did not require support at the Last Assessment; 1 patient was receiving supplemental oxygen at Year 4, and 1 was receiving CPAP at Month 6. Three patients developed the need for respiratory support after Baseline but were weaned before Last Assessment (by Month 9, Year 1.5, and Year 2.5). Of the 24/69 (35%) patients who did require respiratory

support at Baseline (including invasive mechanical ventilation, CPAP, or supplemental oxygen), 11 (46%) no longer required support at Last Assessment.

Growth: Length/height and weight Z-scores generally improved over time (**Figure 4**). Change from Baseline at Last Assessment was significant for both length/height (0.5 [-4, 4]; n=66; $P=.0025$) and weight (1.0 [-5, 6]; n=67; $P=.0001$) Z-scores. Baseline head circumference Z-score was -1.0 (-4, 4; n=56); change from Baseline was 0.1 (-2, 3; n=47) at Month 6 and 0.2 (-3, 7; n=55) at Last Assessment.

Ventilator-free and overall survival: Thirty-eight of the 45 patients (84%) who were not receiving respiratory support at Baseline remained ventilator-free. The Kaplan-Meier estimate of the ventilator-free survival rate at Year 6 for these patients was 84%. Among all 69 patients, the Kaplan-Meier estimate of the overall survival rate at Year 6 was 80%. Survival outcomes for patients in this study were also included in a published analysis that pooled these data with those of a separate study (14).

Other measures

Median (min, max) ALP activity increased from 20 (18, 122) U/L at Baseline (n=65) to 3761 (272, 11,910) U/L after 3 weeks of treatment (n=61) and continued to increase through Year 1 (6742 [1315, 20,041] U/L; n=49). ALP remained elevated throughout treatment, as expected with asfotase alfa treatment. Median (min, max) PPI concentration, which was elevated at Baseline (6.3 [2.7, 13.3] μM ; n=65), decreased to within reference range (1.3–5.7 μM) at Week 6 (3.9 [0.8, 39.2] μM) and remained within

reference range throughout the study. Similarly, median (min, max) PLP concentration decreased from Baseline (521 [48, 24,600] ng/mL; n=60) to within reference range (11.8–68.4 ng/mL) at Week 6 (44 [6, 4590] ng/mL) and remained within reference range through Year 5. PTH levels were 1.2 (0.6, 6.7; n=48) pmol/L at Baseline, 1.7 (0.6, 45.9; n=52) pmol/L at Month 6, and 2.2 (0.6, 10.1; n=66) pmol/L at Last Assessment.

Sixty-two patients had *ALPL* mutation analysis results; 44 patients were compound heterozygous for 2 pathogenic mutations, 9 were homozygous for the same mutant allele, and 9 had only 1 mutation identified consistent with a dominant-negative effect.

Safety and tolerability

All patients experienced ≥ 1 treatment-emergent AE (TEAE). Table 2 summarizes the most common TEAEs occurring in $\geq 20\%$ of patients, regardless of relationship to study drug. Most TEAEs were mild (2125/3052 [70%]) or moderate (728/3052 [24%]) in severity and assessed by the investigator as unrelated to study drug (2409/3052 [79%]).

The most common TEAEs assessed as related to study drug were ISRs (593/643 [92%]) and IARs (11/643 [2%]), which occurred in 43 and 6 patients, respectively. The most common ISRs were injection site erythema (33/69 [48%]), discoloration (12/69 [17%]), induration (11/69 [16%]), and hematoma (10/69 [15%]). IARs consisted of pyrexia (4/69 [6%]), chills (1/69 [1%]), injection site rash (1/69 [1%]), anaphylactoid reaction (1/69 [1%]), drug hypersensitivity (1/69 [1%]), and papular rash (1/69 [1%]).

The IARs of anaphylactoid reaction (categorized as stage 1 anaphylactic shock) and drug hypersensitivity were considered serious; neither patient had received

pretreatment with medications to manage IARs. Both events resulted in interruption of asfotase alfa administration; the anaphylactoid reaction was treated with an IV electrolyte solution, and no treatment was given for the event of drug hypersensitivity. Both events resolved, and treatment with asfotase alfa was restarted without further occurrences.

Lipodystrophy was reported in 5/69 (7%) patients and was mild or moderate in severity and assessed as probably related or related to study drug. Eight patients (8/69 [12%]) had ectopic calcification findings on eye examination, which were identified as TEAEs in 2 patients. Both events involved corneal deposits, were considered unrelated to study drug and did not interfere with vision, and resolved at Last Assessment.

Nephrocalcinosis was reported in 46/69 (67%) patients and was present at Baseline in all but 6 patients. Five patients had nephrocalcinosis reported as a TEAE. An additional 7 had 8 TEAEs that were not recorded as ectopic calcifications or nephrocalcinosis but were considered as such upon medical review. Renal function remained normal in all patients.

A total of 28/69 (41%) patients experienced 46 AEs relevant to the AE of craniosynostosis (onset after start of treatment: 1–1851 d); all but 3 events were assessed as unlikely related or unrelated to study drug, and all but 7 were mild or moderate in severity. Two patients required surgical treatment.

Twenty-two events related to chronic hepatitis were reported in 13/69 (19%) patients. All were mild or moderate in severity. Hepatomegaly in 1 patient was assessed as possibly

related to study drug treatment. A serious AE of increased hepatic enzymes in another patient was moderate in severity and assessed as unlikely related to study drug.

A total of 50/69 (72%) patients experienced 297 serious AEs, most of which (286 [96%]) were assessed by the investigator as unlikely related or unrelated to study drug. Of the 11 serious AEs considered treatment related, 7 were ISRs or IARs in 3 patients; the remaining 4 occurred in 3 patients: craniosynostosis (n=1), pneumonia resulting in study drug withdrawal (n=1), and Arnold-Chiari type 1 malformation and syringomyelia (n=1).

In total, 9 patients (13%) died. The causes of death in 6 patients were respiratory failure and cerebral death (following findings of hypoxia-induced lesions/encephalopathy 1 week prior to death); HPP-related complications; severe respiratory failure; cardiopulmonary arrest; severe cardiopulmonary insufficiency; and transtentorial and cerebellar tonsillar herniation due to cerebral edema related to severe HPP. Three patients died of pneumonia; in 1 patient, pneumonia was considered possibly related to asfotase alfa treatment.

Calcium and magnesium levels were of particular interest in this study population based on their role in bone formation, strength, and rigidity. The mean calcium level was within normal limits at Baseline (mean [SD]: 2.6 [0.3] mmol/L), and only small fluctuations were observed over the course of the study. Post-Baseline changes in calcium levels were considered clinically significant by the investigator in 6 patients; 3 patients had elevated levels (3.6 mmol/L at Week 3 [n=1]; 3.6 mmol/L at Week 3 and 3.3 mmol/L at Week 6

[n=1]; and 3.1 mmol/L at Week 120 [n=1]), of which 2 had elevated levels at Baseline, and 3 had decreased levels (2.1 mmol/L at Month 3 [n=1; age at Baseline: 217 wk]; 2.0 mmol/L at Month 9 [n=1; age at Baseline: 21 wk]; and 2.2 mmol/L at Year 1 [n=1; age at Baseline: 4 wk]). Magnesium levels remained generally within normal range; 1 patient had a clinically significantly low level at Month 3 (0.5 mmol/L) that was normalized at Months 6 and 9 (Last Assessment).

Twelve patients had clinically significantly abnormal hematology findings. Five had low hematocrit and hemoglobin levels, 5 had high leukocyte or lymphocyte counts, 1 had low neutrophil count, and 1 had high blasts. For most patients, hematology findings returned to normal or were no longer considered clinically significant at the patient's Last Assessment; 1 patient had clinically significantly low hematocrit, hemoglobin, and erythrocyte levels at Years 3.5 and 4 (Last Assessment).

Anti-asfotase alfa antibody levels

In total, 60/68 (88%) patients tested positive for anti-asfotase alfa antibodies during the study (maximum titer: 2048); 40 (67%) of these patients tested positive for neutralizing antibodies (NAbs). Six patients tested positive for anti-asfotase alfa antibodies at Baseline, 1 of whom tested negative at all subsequent assessments. The median (min, max) positive NAb titer at Last Assessment, measured as percent inhibition, was 7.7% (4.5, 92.6). Median (min, max) time to detection of first post-Baseline NAb titer was 168.5 (20, 1359) days. No clear relationship was found between the presence of anti-

asfotase alfa antibodies and AEs, nor were any AEs suggestive of immune mediation or tachyphylaxis.

RGI-C “Responders”

Results of the post hoc comparison between RGI-C “responders” (individual mean $\geq +2$ at Year 1) and “nonresponders” at Week 48 are summarized in **Table 3**. Of the 69 patients in this study, 50 (72.5%) of patients had an RGI-C score of $\geq +2$ at Year 1. Nineteen patients did not achieve an RGI-C score of $\geq +2$ at Year 1; median RGI-C score for these “nonresponders” was 0.67 at Year 1. Seventeen of these “nonresponders” had a last overall on-treatment assessment, and of these, 5 achieved a RGI-C score of $\geq +2$ by approximately 2.3 years. In this same timeframe, an additional 4 “nonresponders” achieved a RGI-C score between $+1$ and $< +2$. The remaining 8 “nonresponders” had RGI-C scores $< +1$ at Last Assessment.

A greater proportion of “responders” than “nonresponders” completed the study (94% vs. 68%, respectively; $P=.0105$), with fewer deaths (4% vs. 37%, respectively; $P=.0012$). Compared with “responders,” “nonresponders” also had higher serum calcium ($P=.0204$), plasma PLP ($P=.0403$), and plasma PPI concentrations ($P=.0427$) at Baseline. “Nonresponders” had lower length/height Z-scores ($P=.0345$) and smaller chest circumferences ($P=.0261$) at Baseline than “responders.” Correlation between Baseline ALP activity and RGI-C score at Year 1 was moderate in “nonresponders” (Pearson correlation coefficient: 0.5468; $P=.0189$) and weak in “responders” (0.2008; $P=.1759$). There was no statistical difference between “responders” and

“nonresponders” in anti-asfotase alfa antibody status during the study. Positive NAb status was not statistically significantly different between “responders” and “nonresponders” at Month 6 or at Year 1, but the percentage of “nonresponders” positive for NAb was significantly higher (9/19; 60%) than the percentage of “responders” (9/50; 20%; $P=.0047$) at Last Assessment.

DISCUSSION

This open-label study of asfotase alfa treatment in infants and young children with a follow-up period of up to 6 years is the largest, prospective clinical study of HPP to date. The enrolled population of 69 children with severe HPP is exceptionally large for a study of a rare disease such as HPP. This study assessed therapy response, showing that asfotase alfa was efficacious and safe in the majority of children who had onset of severe HPP signs/symptoms before age 6 months. Improvements in skeletal manifestations, respiratory support, and growth were observed during treatment with asfotase alfa within 6 months and were sustained for a median of 2.3 years and up to 6 years of treatment. Radiographic “nonresponders” at Year 1 generally had more severe disease at Baseline, and a higher incidence of positive NAb status at study end than “responders.”

The primary efficacy measures at Month 6 and Year 1 were met, showing significant improvements in HPP-related skeletal manifestations. Moreover, 58% of patients were considered “responders” at Month 6 and 72% at Year 1, indicating substantial healing of skeletal manifestations. Improvements in respiratory status were also observed; nearly

half (46%) the patients who started the study requiring respiratory support were able to forego support by Last Assessment, and 89% who were free of respiratory support at Baseline remained so over the study. These results are consistent with those of previous smaller studies in infants and young children (n=11; age: 2 wk–3 y) and in Japanese patients with HPP (n=13; median [min, max] age at Baseline: 91 d [0 d, 34 y]) (17,18,23).

Nine deaths occurred in this study, mostly attributed to underlying HPP disease. The majority of deaths (78%) were radiographic “nonresponders.” One death, which was attributed to pneumonia, was considered by the investigator to be possibly related to asfotase alfa. In a prior survival analysis, which included data from some patients in the current study and other asfotase alfa studies, treatment significantly ($P<.0001$) improved survival among patients with perinatal and infantile HPP; survival was 95% at age 1 year and 84% at age 5 years but only 42% and 27%, respectively, among historical controls (14).

Asfotase alfa was generally well tolerated, with an overall safety profile consistent with that observed previously (14,17,23). All patients experienced ≥ 1 TEAE; most were mild or moderate in severity and assessed as unrelated to study drug. Less than 10% of patients experienced serious AEs that were considered treatment related. The majority (64%) of treatment-related serious AEs were ISRs or IARs. Clinicians should be aware of strategies to minimize or prevent ISRs, such as proper injection technique, as well as management options for reactions requiring treatment (24). Post-Baseline elevated

calcium levels were considered clinically significant in 3 patients, 2 of whom had clinically significant elevations at Baseline. Three patients had clinically significant hypocalcemia during the study. Previous reports discussed the role of dietary calcium restriction in the management of HPP before asfotase alfa became available, as many patients have a history of hypercalcemia (17,23). The improved skeletal mineralization associated with asfotase alfa increases calcium intake requirements (i.e., hungry bone syndrome). Hence, supplying sufficient dietary calcium and monitoring serum calcium and PTH during initial treatment are essential (24). Monitoring urine calcium in relation to serum calcium and PTH is also important in guiding calcium requirements.

Anti-asfotase alfa antibodies were identified in 88% of patients during this study, and 67% of these patients tested positive for NAb. Development of anti-drug antibodies has been documented in previous clinical studies of asfotase alfa, with no apparent impact on clinical outcomes (17,25). These data reflect small sample sizes (9–12 patients) and a short duration of treatment exposure (up to 1 year). The current study, which includes the largest population to date of asfotase alfa-treated patients, with a median (min, max) treatment duration of 2.3 (0.02, 5.8) years, showed no clear relationship between the presence of these antibodies and AEs. A post hoc analysis suggests that NAb positive status may be associated with slow radiographic response as assessed by the RGI-C at Year 1. Unfortunately, anti-drug antibody testing is not yet commercially available to identify patients who may not be responding to treatment because of the presence of NAb. The development of NAb could be associated with the severity of the disease, as has been observed in other conditions such as infantile-onset Pompe

disease (26), but to date, the limited data available do not allow us to draw firm conclusions. A recent case report found that the development of NABs (% inhibition: 40.4%) in a patient with HPP treated with asfotase alfa for 2.5 years was associated with loss of efficacy and that immune tolerance induction therapy and a 1-month discontinuation of asfotase alfa successfully restored treatment efficacy within 6 months (27). Currently, anti-drug antibody testing is available for patients enrolled in the Global HPP Registry (NCT02306720; EUPAS13514), which will also allow for the collection of data to further understand the impact of NABs on the efficacy and safety of asfotase alfa.

Generally, RGI-C “nonresponders” at Year 1 represent a subgroup of patients with more severe HPP at Baseline, evidenced by narrower chest walls (i.e., smaller chest circumference), greater requirements for mechanical ventilation, and a higher proportion of deaths. Levels of TNSALP substrates (PPi and PLP) and serum calcium were also higher in “nonresponders.” These characteristics should be considered when determining dosing for patients with severe perinatal HPP. It should also be considered that patients with severe disease may have delayed response to treatment or, in some cases, limited response to treatment due to serious preexisting complications of HPP such as severe hypomineralization or lung hypoplasia.

A limitation of this study was the heterogeneity of study patients, as some patients had life-threatening perinatal disease. In addition, treatment was initiated at various ages, making comparisons difficult. Statistical analyses also did not correct for multiplicity or

confounding variables. Lastly, in our post hoc “responder” analysis, although there were some patients who had “minimal” radiographic improvements, as a result of our strict definition of “responders” (RGI-C score $\geq +2$), these patients were included in the “nonresponder” group. Accordingly, 9 patients in the “nonresponder” group, included based on their RGI-C scores at Year 1, had scores $\geq +1$ to $< +3$ at Last Assessment, indicating minimal or substantial healing.

CONCLUSIONS

Most infants and young children with HPP treated with asfotase alfa showed sustained improvements in HPP-related skeletal manifestations, respiratory function, and growth. Asfotase alfa was generally well tolerated. A subgroup of the study patients with very severe disease at Baseline were classified here as radiographic “nonresponders.” These patients should be closely monitored for therapeutic response, with dose adjustments and additional therapeutic measures considered, as necessary.

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626 ***Data sharing***

627 Qualified academic investigators may request participant-level, de-identified clinical data
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632

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LEGENDS

Figure 1. Patient disposition.

^aAll patients who received any asfotase alfa, regardless of whether they were lost to follow-up or dropped out of the study.

^bAll patients from the full analysis set who did not have any major protocol deviations deemed to potentially influence treatment effect.

^c1 patient did not meet the eligibility criteria and had incorrect administration of study drug.

^dConsidered possibly related to study drug treatment in 1 patient.

Figure 2. Median RGI-C scores over time in infants and children with HPP treated with asfotase alfa. Patients with RGI-C scores of $\geq +2$ were classified as “responders.” RGI-C scale: -3 =severe worsening of HPP-related skeletal manifestations; -2 =moderate worsening; -1 =minimal worsening; 0 =no change; $+1$ =minimal healing; $+2$ =substantial healing; $+3$ =complete or near complete healing.

^aMonth 6 and Year 1 values differ from those in the text for the primary efficacy measure because no imputation of missing values was performed.

^bLast Assessment was defined as the latest post-Baseline assessment on treatment (within 5 days after end of treatment) with a nonmissing value for each patient; overall median (min, max) treatment duration was 2.3 (0.02, 5.8) years.

^cAll patients were included in the full analysis population; the decreasing n is due to the number of patients on treatment at end of study or because assessments may not have been done at each time point.

* $P < .01$ based on Wilcoxon signed-rank test comparing median change to 0.

HPP=hypophosphatasia; RGI-C=Radiographic Global Impression of Change.

Figure 3. Median change from Baseline in RSS scores over time in infants and children with HPP treated with asfotase alfa.

^aLast Assessment was defined as the latest post-Baseline assessment on treatment (within 5 days after end of treatment) with a nonmissing value for each patient; overall median (min, max) treatment duration was 2.3 (0.02, 5.8) years.

^bAll patients were included in the full analysis population; the decreasing n is due to number of patients on treatment at the end of study or because assessments may not have been done at each time point.

* $P < .05$ based on Wilcoxon signed-rank test comparing median change to 0.

HPP=hypophosphatasia; RSS=Rickets Severity Scale.

Figure 4. Median (min, max) Z-scores for (A) length/height and (B) weight over time in infants and children with HPP treated with asfotase alfa.

^aLast Assessment was defined as the latest post-Baseline assessment on treatment (within 5 days after end of study treatment) with a nonmissing value for each patient; overall median treatment duration was 2.3 (0.02, 5.8) years.

^bAll patients were included in the full analysis population; the decreasing n is due to the number of patients on treatment at the end of study or because assessments may not have been done at each time point.

* $P < .05$ based on Wilcoxon signed-rank test comparing median change to 0.

779 HPP=hypophosphatasia; SD=standard deviation.

780

781 **Table 1. Baseline Demographic and Clinical Characteristics**

Baseline Characteristic	Enrolled Patients N=69
Age at enrollment, mo, median (min, max)	16.0 (0.3, 72.2)
Sex, n (%)	
Male	33 (48)
Race, n (%)	
White	54 (78)
Asian	7 (10)
Other	3 (4)
Unknown	5 (7)
Age at first signs of HPP, mo	
Median (min, max)	1.0 (0, 5.5)
HPP-specific medical history, n (%)	
Abnormally shaped chest	58 (84)
History of respiratory compromise (up to and including respiratory failure) ^a	46 (67)
Seizures	17 (25)
Difficulty gaining weight, failure to thrive, and/or difficulty eating/swallowing	60 (87)
Hypercalcemia	61 (88)
Nephrocalcinosis	37 (54)
Fractures and/or delayed fracture healing	21 (30)
Length/height Z-score	n=67
Median (min, max)	-2.7 (-10.0, 1.0)

Baseline Characteristic	Enrolled Patients N=69
Weight Z-score	n=68
Median (min, max)	-2.5 (-24.0, -0)
RSS score	n=67
Median (min, max)	4.0 (0.0, 10.0)
ALP, U/L [normal range: 60–370 U/L] ^b	n=65
Median (min, max)	20 (18, 122)
PPI, µM [normal range: 1.3–5.7 µM]	n=65
Median (min, max)	6.3 (2.7, 13.3)
PLP, ng/mL [normal range: 11.8–68.4 ng/mL] ^c	n=60
Median (min, max)	521 (48, 24600)
Calcium, mmol/L [normal ranges: 2.25–2.74 mmol/L (age: ≤2 y); 2.1–2.57 mmol/L (age: >2 y)]	n=65
Median (min, max)	2.6 (1.8, 4.0)

^aRespiratory compromise was defined as respiratory signs/symptoms that required institution of respiratory support measure(s), required medication(s) for management of symptom(s), and/or were associated with other respiratory complications (e.g., pneumonia, respiratory tract infection).

^bNormal range for ALP activity per ARUP Laboratories (University of Utah, Salt Lake City, UT) varies by age: 0–30 days: 60–320 U/L; 1–11 months: 70–350 U/L; 1–3 years: 125–320 U/L; 4–6 years: 150–370 U/L. Normal range also varies by sex in patients older than 10 years of age.

^cMedian (min, max) concentration for patients receiving vitamin B6 supplementation before dosing (n=14) was 9960 (65, 24600) ng/mL and for those patients not receiving vitamin B6 supplementation before dosing (n=46) was 417 (48, 13100) ng/mL.

ALP=alkaline phosphatase; HPP=hypophosphatasia; PPI=inorganic pyrophosphate; PLP=pyridoxal 5'-phosphate.

796 **Table 2. Treatment-Emergent Adverse Events Occurring in >20% of Patients—**
797 **Safety Analysis Set**

TEAE ^a	Patients, n (%) ^b
Pyrexia	47 (68)
Tooth loss	41 (59)
Injection site erythema	33 (48)
Vomiting	31 (45)
Diarrhea	20 (29)
Craniosynostosis	19 (28)
Upper respiratory tract infection	19 (28)
Nasopharyngitis	18 (26)
Gastroenteritis	17 (25)
Cough	17 (25)
Respiratory tract infection	16 (23)
Constipation	16 (23)
Pneumonia	14 (20)

798 ^aAdverse events coded using MedDRA Version 13.0.
799 ^bPatient percentages are based on the total number of patients in the treatment group
800 (N=69).
801 TEAE=treatment-emergent adverse event.
802

803 **Table 3. Comparison of Baseline Characteristics and Outcomes in Radiographic**
804 **“Responders” vs. “Nonresponders”^{*}—Full Analysis Set**

	“Responders” (n=50)	“Nonresponders” (n=19)	P Value
Disposition			
Completed study, n (%)	47 (94)	13 (68)	.0105
Discontinued, n			
Withdrawal by parents	1	2	
Adverse event	2	4	
Characteristics			
Age at enrollment, mo, median (min, max)	21.0 (0, 71.4)	8.9 (0.4, 71.7)	.2318
Male, n (%)	25 (50)	8 (42)	.5997
Age at HPP onset, mo, median (min, max)	1.0 (0, 5.5)	1.0 (0, 5.0)	.1164
Time from HPP diagnosis to treatment, mo, median (min, max)	20.3 (0, 67.8)	8.1 (0.3, 67.3)	.3753
Weight Z-score	n=49	n=19	
Median (min, max)	-2.3 (-7.8, -0.04)	-2.7 (-23.8, -0.3)	.3930
Length Z-score	n=48	n=19	
Median (min, max)	-2.6 (-8.3, 0.9)	-3.5 (-10.1, -0.3)	.0345
Chest circumference, cm	n=47	n=19	
Median (min, max)	42.5 (32.0, 56.0)	37.0 (27.5, 51.5)	.0261
Race			
Asian, n (%)	5 (10.0)	2 (10.5)	1.0000
White, n (%)	38 (76.0)	16 (84.2)	
Multiple, n (%)	1 (2.0)	0	
Other, n (%)	2 (4.0)	0	
Unknown/not reported, n (%)	4 (8.0)	1 (5.3)	

	“Responders” (n=50)	“Nonresponders” (n=19)	P Value
Laboratory Parameters, median (min, max)			
ALP, U/L	n=47 23 (18, 122)	n=18 18 (18, 50)	.0513
Calcium, mmol/L	n=47 2.5 (1.8, 3.6)	n=18 2.6 (2.3, 4.0)	.0204
PLP, ng/mL	n=43 429.0 (65.1, 24100.0)	n=17 1300.0 (47.5, 24600.0)	.0403
PPi, μ M	n=46 5.9 (2.7, 12.5)	n=19 7.1 (3.6, 13.3)	.0427
Magnesium, mmol/L	n=47 0.9 (0.6, 1.2)	n=18 0.9 (0.6, 1.1)	.6433
Phosphate, mmol/L	n=47 2.0 (1.2, 2.5)	n=18 2.0 (0.9, 2.7)	.7918
Parathyroid hormone, pmol/L	n=34 1.5 (0.6, 5.4)	n=14 0.6 (0.6, 6.7)	.7302
ALPL Gene Polymorphism, n (%)	n=45	n=17	.6634
Compound heterozygous	31 (69)	13 (77)	
Heterozygous	8 (18)	1 (6)	
Homozygous	6 (13)	3 (18)	
Disease Characteristics, n (%)			
Respiratory compromise	33 (66)	12 (63)	1.000
Rachitic chest	49 (98)	17 (90)	.1817
B ₆ -responsive seizures	11 (22)	8 (42)	.1320
Abnormally shaped chest	39 (78)	13 (68)	.5327
High serum calcium	35 (70)	14 (74)	1.000
Nephrocalcinosis	25 (50)	12 (63)	.4208
Mean baseline RSS (SD)	4.7 (3.1)	4.8 (3.5)	.9833

	“Responders” (n=50)	“Nonresponders” (n=19)	P Value
Respiratory Characteristics			
Fraction of inspired oxygen, %	n=12	n=7	
Median (min, max)	29.0 (21.0, 50.0)	25.0 (21.0, 54.0)	.3500
Inspiratory pressure, cm H ₂ O	n=8	n=5	
Median (min, max)	25.5 (0.4, 31.0)	24.0 (15.0, 32.0)	.6601
Expiratory pressure, cm H ₂ O	n=9	n=7	
Median (min, max)	7.0 (5.0, 12.0)	6.0 (5.0, 8.0)	.2726
Respiratory support type, n (%)			
No support	35 (70)	10 (53)	.5079
Mechanical ventilation	8 (16)	5 (26)	
Supplemental oxygen	4 (8)	2 (11)	
CPAP	2 (4)	2 (11)	
Other	1 (2)	0 (0)	
Respiratory support duration, h	n=14	n=8	
Median (min, max)	24 (8, 24)	24 (24, 24)	.4497
Treatment Exposure			
Dosing frequency, n (%)			.6642
Always received 3 times/wk	46 (92)	17 (90)	
Ever received >3 times/wk	4 (8)	2 (11)	
Last dose received, mg/kg/wk, median (min, max)	4.0 (2, 12)	5.7 (2, 11)	.2924
Treatment duration, y, median (SD)	2.3 (0.6, 8.8)	1.9 (0.02, 4.2)	.1342
ADA/NAbs			
Positive ADA status, n (%)			
Month 6	34 (71)	12 (80)	.7400
Year 1	28 (64)	9 (90)	.1411
Ever	45 (90)	15 (83)	.4279
Positive NAb status, n (%)			
Month 6	10 (30)	6 (50)	.2963

	“Responders” (n=50)	“Nonresponders” (n=19)	P Value
Year 1	8 (29)	5 (56)	.2293
Ever	29 (64)	11 (73)	.7529
Last Assessment	9 (20)	9 (60)	.0074
NAb by percent inhibition, median (min, max)			
Month 6	2.4 (−1.7, 90.3)	5.4 (−0.5, 88.5)	.1820
Year 1	2.5 (−6.2, 63.5)	4.7 (−3.5, 93.8)	.3046
Death			
Number of deaths on study, n (%)	2 (4)	7 (37)	.0012
RGI-C Scores			
RGI-C score at Last Assessment	n=50	n=17	
Median (min, max)	+2.7 (−2.7, +3.0)	+1.0 (−1.7, +2.7)	<.0001
Category, n (%)			
−3 to <−2	2 (4)	0	
−2 to <−1	0	1 (6)	
−1 to <0	1 (2)	4 (24)	
0 to <+1	0	3 (18)	
+1 to <+2	3 (6)	4 (24)	
+2 to <+3	40 (80)	5 (29)	
+3	4 (8)	0	

ADA=anti-drug antibodies; ALP=alkaline phosphatase; CPAP=continuous positive airway pressure; HPP=hypophosphatasia; NAb=neutralizing antibody; RGI-C= Radiographic Global Impression of Change; PLP=pyridoxal 5'-phosphate; PPi=inorganic pyrophosphate.

*RGI-C “responders”: mean score $\geq +2$ at Year 1.

Figure 1.

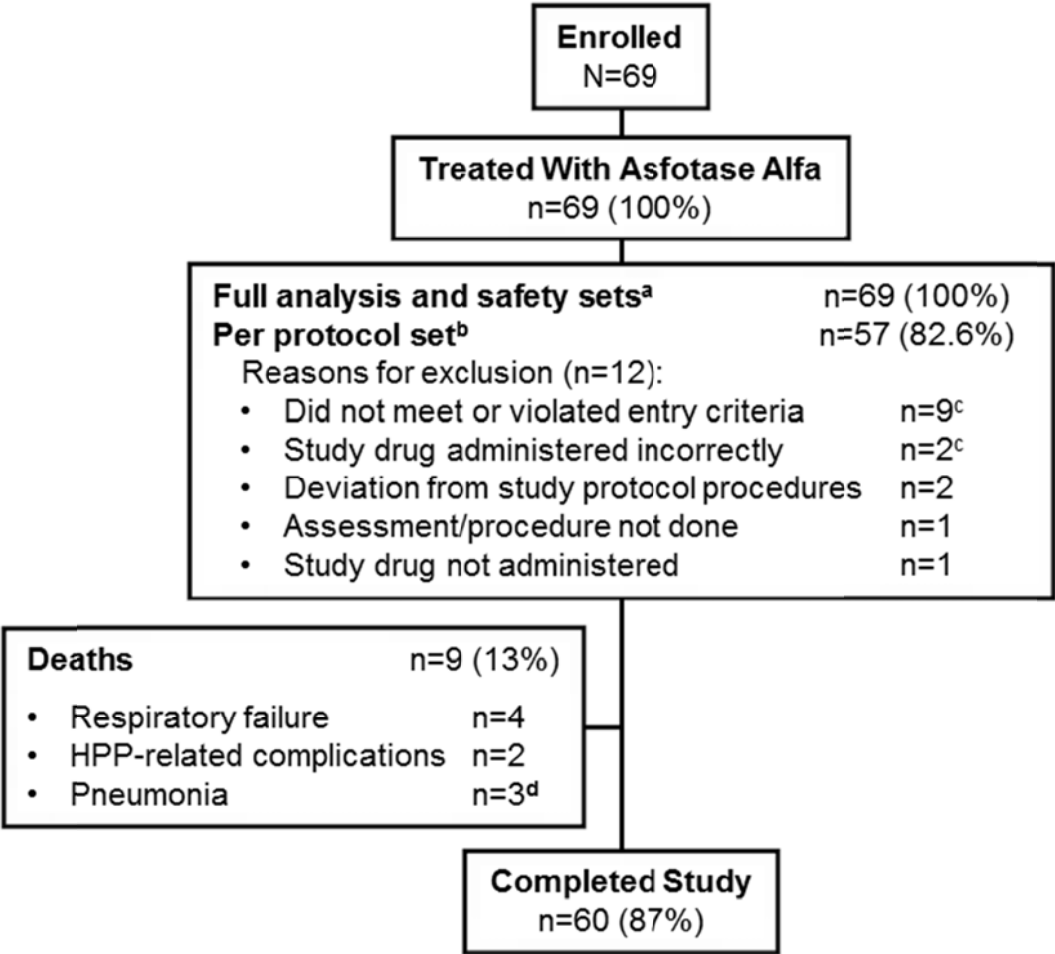


Figure 2

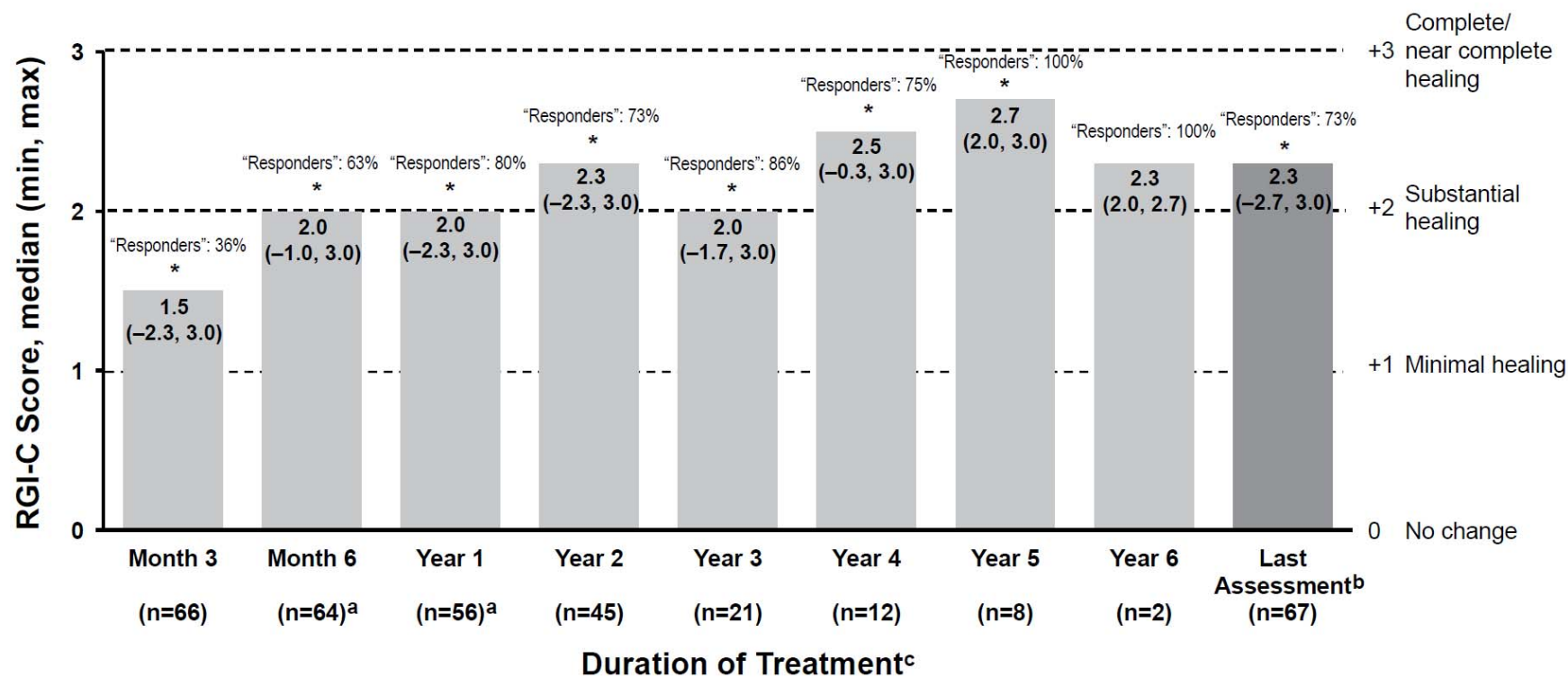


Figure 3

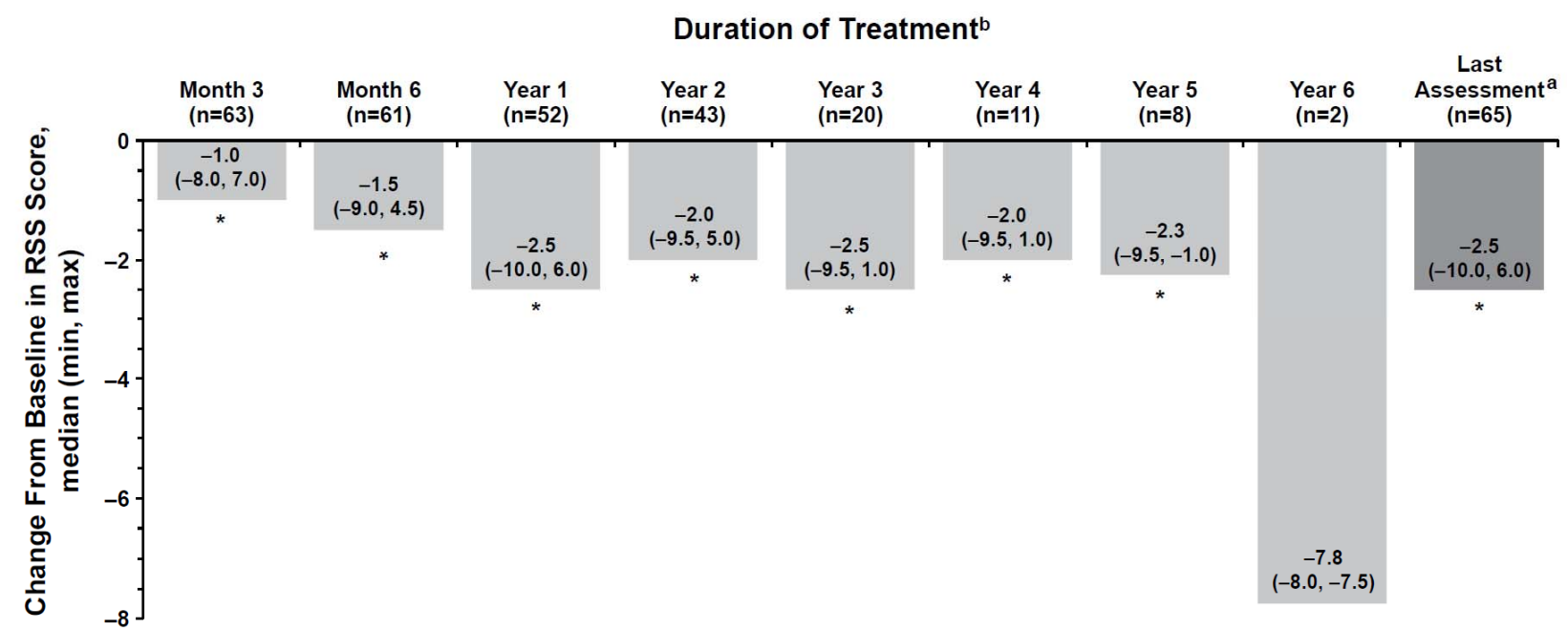
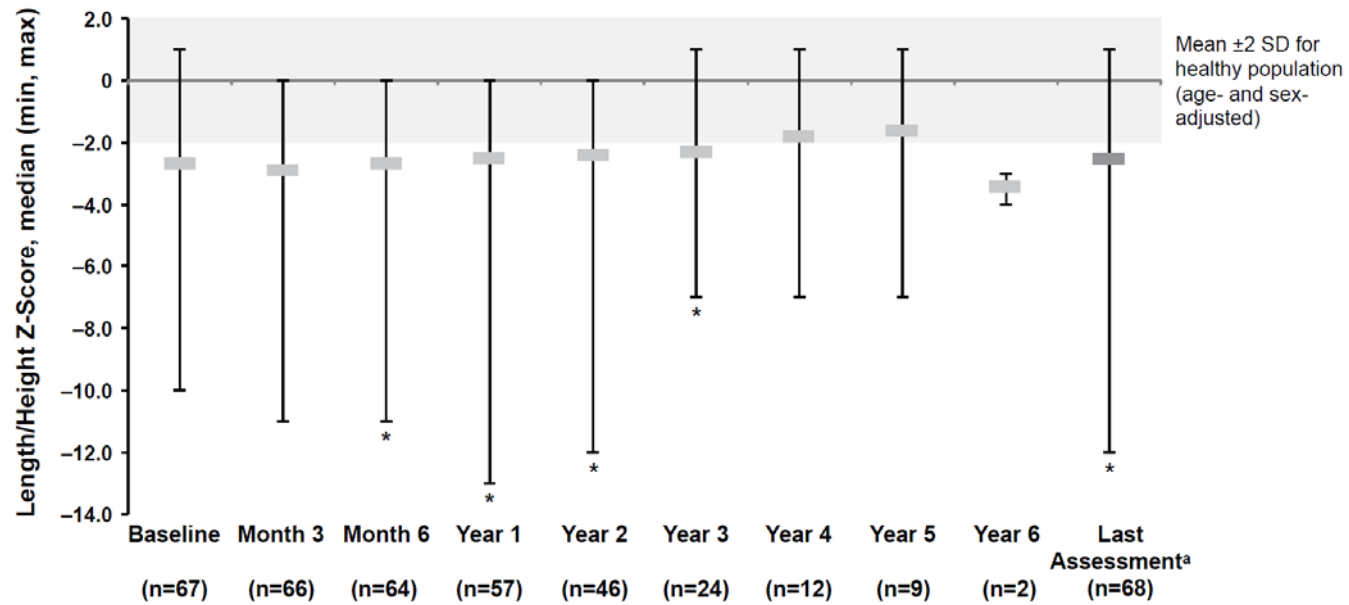


Figure 4

A Length/Height



B Weight

